PTO/SB/08a (11-07) owed for use through 11/30/2007 OMB 0651-0031

Under the	U Paperwork Roduction Act of 1995, no cersons are require	IS Patient and Trademark Office; U.S. DEPARTMENT OF COMMERCE of tomecond to a collection of information unless it contains a valid OMB control number	
Substitute for form 1449A/PTO	Complete if Known		
	Application Number	10/575,132	
INFORMATION DISCLOSURE	Filing Date	July 7, 2006	
STATEMENT BY APPLICANT	First Named Inventor	Sarah Donald	
	Art Unit	1612	
(Use as many sheets as necessary)	Examiner Name	Chris E. Simmons	
Sheet 1 of 3	Attorney Docket Number	13566.105014	

			U.S	. PATENT D	ОСИМЕ	NTS		
Examiner Cite		Document Number	Publication Date Name of Patentee of		Patentee or Applicant of Cited Document	r Applicant of Pages, Columns, Lines, Where Releva		
Initials *	No.1	Number - Kind Code <sup>2</sup> (If known)	м	MM-DD-YYYY		ONLY DACOHER	Passages or Relevant Figures Appear	
		F	ORE	IGN PATENT	T DOCU	MENTS		
Examiner	Cite	Foreign Patent Document			Name of Patentee or		Pages, Columns, Lines, Where Relevant	
Initials*		Publication MM-DD-Y		Applicant of Cited Document	Passages or Relevant Figures Appear	T°		
/C.S./		WO 03/020259	)	03-13-2	2003	Cancer Research Technology Limited		
	_							
								_
				-				-

Signature	/Onna Onnanona/	Considered		
*EXAMINER Initial if r	reference considered, whether or not citation is in conformance with MPEP 609	Draw line through citation if n	of in conformance and not considered. Include copy of this form with next	1
communication to appl	ficant "Applicant's unique citation designation number (optional) 2 See Kinds	Codes of USPTO Patent Door	aments at <a href="https://www.usplo.gov">www.usplo.gov</a> or MPEP 901 04 * Enter Office that issued the	,
	etter code (WIPO Standard ST 3). 4 For Japanese patent documents, the indicate			1
document by the appro	priate symbols as indicated on the document under WIPO Standard ST, 16 if pos	sible." Applicant is to place a ch	nock mark here if English language Translation is attached	
This collection of infor	mation is required by 37 CFR 197 and 198. The information is required to	obtain or retain a benefit by the	he public which is to file (and by the USPTO to process) an application.	

Date

05/17/2009

USPTO Time will vary depending upon the individual case. Any comments on the amount of time you require to cominformation Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA.2.

/Chric Simmone/

Examiner

1	Substitute	for form 1449A/PTC	)			Complete if Known
ı					Application Number	10/575,132
ı	INFO	RMATION	DIS	CLOSURE	Filing Date	July 7, 2006
ı	STATEMENT BY APPLICANT				First Named Inventor	Sarah Donald
ı					Art Unit	1612
L		(Use as many she	ets as	necessary)	Examiner Name	Chris E. Simmons
$\overline{}$	Sheet	2	of	3	Attorney Docket Number	13566.105014

		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials *	Cite No.1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, oty) and/or country where published.	T²
		Alexopoulos, "Phase II study of pegylated liposomal doxorubicin (Caelyx(R)) and docetaxel as first-line treatment in metastatic breast cancer," Ann. Oncol., 2004, 15(6):891-5	
		D'Incalci et al., "Unique Features of the Mode of Action of ET-743", The Oncologist, 7, p. 210-216, June 2002	
		Donald et al, "Comparison of four modulators of drug metabolism as protectants against the hepatotoxicity of the novel antitumor drug yondelis (ET-743) in the female rat and in hepatocytes in vitro," Cancer Chemother Pharmacol, April 2004, vol. 53, pp. 305-12	
		European Medicines Agency (EMEA), "Scientific Discussion" from the European Public Assessment Report for Yondelis®, Revision 1, published March 31, 2008, downloaded from the internet on April 2, 2008, from the website < <a href="http://www.emea.europa.eu/humandocs/Humans/EPAR/yondelis/yondelis.htm">http://www.emea.europa.eu/humandocs/Humans/EPAR/yondelis/yondelis.htm</a> >	
		Forouzesh et al., Proc. Am. Soc. Clin. Oncol. ASCO meeting, Abstract 373, June 3, 2001, Internet Archive Entry from the website < <a "="" href="http://wwb.archive.org/web/" http:="" www.asco.org="">, 32 pages</a>	
		Gourley C. et al., "Malignant mixed Mesodermal Tumours - Biology and Clinical Aspects," European Journal of Cancer, 2002, vol. 38, no. 11, pages 1437-1446	
		Halm et al., "A phase II study of pegylated liposomal doxorubicin for treatment of advanced hepatocellular carcinoma," Ann. Oncol., 2000, 11(1):113-114	
		Hoekman at al., "A phase I/II study of dose-escalated docetaxel given two weekly in combination with a fixed dose of G-CSF," European Journal of Cancer, vol. 37, page \$76, Abstract 270, October 22, 2001	

/Chris Simmons/

05/17/2009

							_
1	Substitute for form 1449A/PTO				ĺ	Complete if Known	
ı					Application Number	10/575,132	
ı	INFO	RMATION	DIS	CLOSURE	Filing Date	July 7, 2006	
ı	STATEMENT BY APPLICANT				First Named Inventor	Sarah Donald	
ı					Art Unit	1612	
L		(Use as many she	ets as	necessary)	Examiner Name	Chris E. Simmons	
١	Sheet	3	of	3	Attorney Docket Number	13566.105014	

		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials *	Cite No.1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, (iy) and/or country where published.	T <sup>2</sup>
		Horstmann et al., "Risks and Benefits of Phase I Oncology Trials, 1991 though 2002," New England Journal of Medicine, vol. 352, pages 895-904; March 3, 2005	
		Lau et al., "A Phase I and Pharmacokinetic Study of Ecteinascidin-743 (Yondelis) in Children with Refractory Solid Tumors." Clinical Cancer Research, vol. 11, pp. 672-677, Jan. 15, 2005	
		PR Newswire, PR Newswire, October 14, 2001, 4 pages	
		Puchalski et al., "Pharmacokinetics of Ecteinascidin 743 Administered as a 24-h Continuous Intravenous Infusion to Adult Patients with Soft Tissue Sarcomas associations with Clinical Characteristics, Pathophysiological Variables and Toxicity," Cancer Chemotherapy and Pharmacology, 2002, vol. 50, no. 4, pages 309-319	
		Rote Liete 2002 "Dexerubicin," entrice 86 056 through 86 062, 2002 not in English	
		Sarosy et al., "Phase I Study of α2-interferon plus Doxorubicin in Patients with Solid Tumors," Cancer Research, vol. 46, pp. 5368-5371, 1986	
		Schwartsmann G. et al., "Marine Organisms as a Source of New Anticancer Agents," The Lancet Oncology, 2001, vol. 2, no. 4, pages 221-225	
		Twelves et al., "Phase I and pharmacokinetic study of YondelisTM (Ecteinascidin-743; ET-743) administered as an infusion over 1 h or 3 h every 21 days in patients with solid tumours," European Journal of Cancer, vol. 39, p. 1842-1851, 2003; available online August 14, 2003	
		Wollina, "Multicenter study of pegylated liposomal doxorubicin in patients with cutaneous T-cell lymphoma," Cancer 2003, 1:98(5):993-1001, published online July 24, 2003	

Examiner Signature	/Chris Simmons/	Date Considered	05/17/2009
-----------------------	-----------------	--------------------	------------

<sup>\*</sup>EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered.

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

<sup>\*</sup>EXAMINES: Indial if reference considered, whether or not classion is in conformance with MPEP 000. Dawn line through classion if not in conformance and not considered.

\*\*Applicant's using a classification designation. Amplicant is bytical as check white her English impaging Translation is established.

\*\*This collection of information is required by 37 CHR 1.98. The information is required to closing or invalidation of religional consideration. Confidence of the confidence